FDA 510(k) Premarket Notification

K990665 BQ/BM INTEGRATED LASER DELIVERY SYSTEM

CLEARED

May 17, 1999
76 days to decision

K990665 · Product code: **HQF** · Ophthalmic
Source: https://www.510kdatabase.net/k990665/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Laser, Ophthalmic (HQF)
Date received	Mar 2, 1999
Decision date	May 17, 1999
Days to decision	76 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**American Laser Corp.**
Location	Mchenry, IL, US
Contact	DANIEL HOEFER
510(k) history	5 submissions · 5 cleared · 1982-2001